AMENDMENTS TO THE CLAIMS:

- 1-36. (Canceled).
- 37. (Currently amended) A drug delivery matrix, comprising a copolymer of ethylene with carboxylic acid and a drug contained within or attached to the matrix, wherein the copolymer is a coating on an implantable substrate.
- 38. (Previously presented) The drug delivery matrix of claim 37, wherein the carboxylic acid co-monomer content is no less than 5% by weight.
- 39. (Previously presented) The drug delivery matrix of claim 38, wherein the carboxylic acid co-monomer content is no more than 50% by weight.
- 40. (Previously presented) The drug delivery matrix of claim 37, wherein the carboxylic acid co-monomer content is no more than 50% by weight.
- 41. (Previously presented) The drug delivery matrix of claim 37, wherein the copolymer is ethylene acrylic acid.
- 42. (Previously presented) The drug delivery matrix of claim 37, wherein the carboxylic acid is selected from a group consisting of acrylic acid, methacrylic acid, maleic acid, itocanic acid, and esters thereof.
- 43. (Canceled).
- 44. (Allowed) A method of coating an implantable medical device, comprising:

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adding a copolymer of ethylene with carboxylic acid to a solvent system to form a composition;

applying the composition to an implantable medical device; and allowing the solvent system to evaporate.

- 45. (Allowed) The method of claim 44, wherein the carboxylic acid is selected from a group consisting of acrylic acid, methacrylic acid, maleic acid, itocanic acid, and esters thereof.
- 46. (Currently amended) The method of claim 44, wherein adding the copolymer to the solvent system further comprises neutralizing the copolymer in a volatile or a non-volatile base and dispersing the copolymer in water and/or a co-solvent eosolvents.
- 47. (Allowed) The method of claim 44, further comprising adding a therapeutic agent to the solvent system.
- 48. (Allowed) The method of claim 44, wherein the solvent system comprises toluene.
- 49. (Allowed) The method of claim 48, wherein the solvent system further comprises a chlorinated solvent and a lower alcohol.
- 50. (New) The method of claim 44, wherein the carboxylic acid co-monomer content is no less than 5% by weight.
- 51. (New) The method of claim 50, wherein the carboxylic acid co-monomer content

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is no more than 50% by weight.

- 52. (New) The method of claim 44, wherein the carboxylic acid co-monomer content is no more than 50% by weight.
- 53. (New) The method of claim 44, wherein the co-polymer is ethylene acrylic acid.
- 54. (New) The method of claim 44, wherein the device comprises a stent.
- 55. (New) The drug delivery matrix of claim 37, wherein the implantable substrate comprises at least a portion of a stent body.